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(71) Applicant and

(72) Inventor: FOREMAN, Paul, David [GB/GB]; 57 Herbert Road, Rainham, Kent ME8 9DA (GB).

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(72) Inventor: ZAREBSKI, Andrew; c/o Primasil Silicones Limited, Kington Road, Weobley, Herefordshire HR4 8QU (GB).

(74) Agents: TOLLETT, Ian et al.; Williams Powell, Morley House, 26-30 Holborn Viaduct, London EC1A 2BP (GB). (81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.

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(54) Title: GLAZING JOINT



(57) Abstract: A composition for glazing joints comprises (a) a siloxane polymer having a molecular weight of from 300,000 to 700,000, (b) a siloxane polymer having a molecular weight of from 10,000 to 100,000 (the ratio of component A to component B being in the range of from 10 to 1, to 3 to 1) and a cross linking agent. The composition is preferably sufficiently cross-linked to have an effective surface tackiness. A glazing strip (10) shaped for receiving two or more sheets may be formed from the composition. An advantage of the composition is that it remains adhesive so that in use, this allows glass sheets to be repositioned and removed when dismantling a partition wall.



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GLAZING JOINT

The present invention relates to a composition for use as a glazing joint, especially for glazed partition walls.

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Glazed partition walls are formed by joining a series of frameless glass sheets. Adjacent edges of the glass sheets are typically joined by coating the edge of a first glass sheet with a sealant, such as silicone or mastic, then positioning the edge of a second glass sheet against the coated edge of the first sheet. A problem with this method is that excess sealant is pushed out of the joint and onto adjacent surfaces of the glass sheets and has to be removed, for example by wiping whilst the sealant is still fluid. This results in an inconsistency in the quality of the finish of the joints. Furthermore, gaining access to both sides of the glazed partition wall in order to remove excess sealant can prove difficult.

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UK patent 2,322,663 describes a glazing joint in which an adhesive strip such as double sided adhesive tape, is attached to one side of the joint to stop the silicone from spilling out on that side. Excess silicone pushed out from the other side of the joint still needs to be wiped away.

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UK patent 2,339,589 describes a reusable joint for joining panes of glass. The joint is a polycarbonate strip. Different designs of strips permit joints between 2, 3 or 4 panes of

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glass. The strip has resilient fins which retain the glass sheets. As no adhesive is used, the panes of glass can be accidentally pulled out of the joint.

US patent 6,194,508 (Wacker-Chemie) describes a peroxidically crosslinkable silicone rubber composition for jointing compounds which has reduced yellowing. The composition comprises an organosiloxane, a silicone resin and an organic peroxide.

US patent 4,797,439 (Rhone Poulenc Chemie) and US 5,833,798 (Dow Corning) describe liquid glazing sealants which are curable into an adhesive state in the presence of moisture.

The present invention seeks to provide an improved glazing joint and to address the above disadvantages.

- According to a first aspect of the invention there is provided a composition for glazing joints comprising
 - (a) a siloxane polymer having a molecular weight of from 300,000 to 700,000;
 - (b) a siloxane polymer having a molecular weight of from 10,000 to 100,000; the ratio of component (a) to component (b) preferably being in the range of from 10 to 1, to 3 to 1; and
 - (c) a cross linking agent.

The composition is preferably sufficiently cross-linked to have an effective surface tackiness.

An advantage of the compound is that it remains adhesive (tacky). In use, this allows glass

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sheets to be repositioned and removed when dismantling a partition wall.

The siloxane may comprise polydimethylsiloxane (PDMS), polyvinylmethylsiloxane, polydiphenylsiloxane, polytrifluoropropylmethylsiloxane or

5 polytrifluorovinylmethylsiloxane.

identity of said components.

In the silicone industry, the generic term for a relatively high molecular weight, relatively viscous siloxane (like component (a)) is a "base" and the term for a relatively low molecular weight, relatively low viscosity siloxane (like component (b)) is a "fluid". These terms will be used for components (a) and (b) throughout the present specification. However, it should be appreciated that this is not to be taken as a limitation to the

Advantageously the compound includes PDMS base and PDMS fluid. PDMS bases and fluids from manufacturers such as Rhodia, Dow and Wacker are suitable.

The combination of base and fluid is sufficiently crosslinked so that the compound has a surface tackiness. The ratio of base to fluid may be in the range of from 10 to 1, to 3 to 1. Preferably the ratio is 4 to 1.

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Advantageously, the PDMS base has a molecular weight in the range of from 300,000 to 700,000. The base preferably has a density of about 1.16 cm⁻³. The Shore hardness of

the base may be in the range of from 10 to 95 °A, preferably 40 to 90 °A. The base preferably includes vinyl groups.

The compound may include from about 50 to about 95% PDMS base, preferably from about 80 to about 90%, and most preferably 86% PDMS base.

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Preferably the PDMS fluid has a molecular weight in the range of from 10,000 to 100,000. The fluid preferably has a density of 0.98 s.g. Advantageously, the fluid is low viscosity (i.e. less than 100,000 centistokes). The fluid has less internal cross linking than the base. The fluid preferably does not contain any vinyl groups.

The compound may include from about 50 to about 95% PDMS fluid, preferably from about 5 to about 20%, and most preferably 13% PDMS fluid.

- Preferably, the cross linking agent is a free radical initiator. The free radical initiator may be an organic peroxide. Instead of a free radical initiator, a platinum catalyst may be used. This would require very few changes to the curing process, but may eliminate the need for a hot box.
- The compound may include from about 1% to about 10%, preferably 5% organic peroxide. The percentages given are percentages by weight. The amount of crosslinking agent is sufficient to result in partial cross linking of the composition so that the

composition has a surface tackiness. The organic peroxide is preferably 2,4 Dichlorobenzylperoxide.

According to a second aspect of the present invention there is provided a glazing strip

5 shaped for receiving two or more sheets wherein the surface of the strip is capable of
remaining adhesive. The glazing strip preferably is formed from the composition as
defined above.

An advantage is that the adhesive nature of the strip firmly secures the glass sheets in

position thus reducing the likelihood that they can accidentally be pulled out of the joint.

No additional adhesive is required and so there is no excess sealant to be wiped up. The strip allows the easy assembly of glass partition walls therefore.

Preferably, the strip has recesses for receiving sheets; the angle of a recess being about 65-90 degrees.

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The glazing strip may be transparent, translucent or opaque. The glazing strip may be coloured for use with coloured glass.

The glazing strip may comprise polyurethane or a thermoplastics material, although these materials suffer from ageing and discolouration and have a worse performance in fire.

The strip may be UV resistant.

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According to a third aspect of the present invention, there is provided a method of manufacturing a compound for glazing joints as claimed comprising

- (a) mixing components (a), (b), and (c) as defined above at a temperature of less than about 40 degrees Centigrade to achieve a substantially homogeneous mixture,
- 5 (b) extruding the mixture,
 - (c) curing the mixture at a temperature of from 100 to 900 degrees Centigrade.

Any polymer mixing technique carried out at a temperature of less than about 40 °C can be used, so long as the result is a homogeneous mixture.

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Preferably, step (c) uses a hot air vulcanising unit. The method may involve an additional curing step of from 100 to 300 degrees Centigrade.

According to a fourth aspect of the present invention, there is provided a method of joining glazing sheets comprising

- (a) providing an adhesive strip shaped for receiving two or more glazing sheets, the surface of the strip being inherently adhesive;
- (b) inserting the strip between adjacent edges of at least a first and second sheet.

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It will be apparent that the glazing strip could be used for joining glass, plastic, perspex, or opaque sheets. The glazing strip could be used in vertical or horizontal joints or joints of other angles. The strip may be used with in conjunction with a contact adhesive.

Preferred embodiments of the present invention are described below, by way of example only, with reference to the accompanying drawings, in which:

Figure 1 is a top plan view of a glazing strip for forming a 180 degree joint;

Figure 2 is a top plan view of a joint formed by the glazing strip in Figure 1;

Figure 3 is a front view of the joint in Figure 2;

Figure 4 is a top plan view of a glazing strip for forming a 45 degree joint;

Figure 5 is a top plan view of a glazing strip for forming a 90 degree joint; and

Figure 6 is a top plan view a glazing strip for forming a joint between three glass sheets.

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Preferred dimensions are given in mm. These can change by up to a factor of 3 or more.

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Referring to Figures 1 to 3, a first embodiment of the glazing strip 10 has recesses 12,14 each for receiving a sheet of glass 16,18. The angle of the recess ("a") is 80 degrees. The strip 10 forms a joint between the glass sheets 16,18 such that the sheets are firmly attached to each other. Typically, the strip 10 maintains a distance of 3 mm between the glass sheets 12,14 (distance "d"). The glazing strip 10 forms a 180 degree joint between the two adjacent sheets of glass 16,18.

The glazing strip 10 is formed by extrusion of a material which is capable of remaining of adhesive. The material is a mixture of polydimethylsiloxane (PDMS) base, (Rhodia MF180^m) which is compounded with inert fillers such as fumed silica, PDMS fluid (Rhodia 47V60000^m) and 2,4 dichlorobenzyl peroxide by combining the above components in a ratio of 86 (PDMS base): 13 (PDMS fluid): 1 (organic peroxide).

Rhodia 47V60000 is a linear dimethylpolysiloxane, which has a low surface tension, good thermal stability and a viscosity of about 60,000 centistokes.

The components are mixed thoroughly in a clean environment using either an open mill or an internal mixer, at a temperature of less than 40 degrees Centigrade to avoid curing, and until a homogenous mass is achieved.

The duration of the mixing is dependant upon the size of the batch and as such varies.

The resultant compound is tested for homogeneity by rheometry and specific gravity testing against a standard.

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After mixing, the material is cured using a hot air vulcanising unit (hot box) which is temperature controlled to within +/- 5 degrees Centigrade. The unit is set to a temperature of between about 200 and about 900 degrees Centigrade depending upon the type of production unit being used to feed extrudate to it. The duration of the curing step depends on the rate of extrusion. The temperature and rate of extrusion will vary according to the cross sectional volume of the profile. The extrusion is passed through the hot box at a rate sufficient to achieve the desired state of cure, such as either a partial (skin) cure or full (cure). In the case of a skin cure extrusion, the product will pass directly from the hot box to a curing tunnel where the extrusion will pass through a heated zone (100 to 295 degrees Centigrade) whilst supported by a moving belt. Certain sizes of extrusion will require only belt cure. The heat source is typically radiant elements but could equally be infra red or microwave.

The material is then further cured at about 200 degrees Centigrade for up to 16 hours.

This polymeric compound is designed to exhibit a certain degree of surface tackiness.

5 Typical physical properties of the material are as follows:

TEST	UNITS	1.21 +/- 0.03		
DENSITY	g/cm³			
HARDNESS	Shore °A	60		
TENSILE STRENGTH	Мра	9.21		
ELONGATION @ BREAK	%	543		
TEAR STRENGTH	N/mm	38		
COMPRESSION SET 24 HRS @ 100°C (Recovery=20mins. in air @ 20°C)	%	23		

The material should have a good blend of physical properties.

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The glazing strips can be made in a variety of sizes and thicknesses for joining glass

sheets of different thickness (typically 6-16mm), for example standard toughened or
laminated glass.

In use, the appropriate sized glazing strip is chosen for the type of glass used and the angle of joint required. A first glass sheet 16 is inserted into the recess 12. The strip 10 is adhesive and so the glass sheet 16 is firmly held in place. A second glass sheet 18 is then positioned in recess 14. The edges 20,22 of the strip 10 are flush with the glass sheets 16,18.

A coating may be applied to one or both edges 20, 22 of the strip 10 by spraying or brushed on; the surface of the coating being non adhesive. A typical coating is sprayed silica. Alternatively, at least one of the edges 20, 22 of the strip may be dried such that at least the outer surface of the strip is no longer sticky.

Referring to Figure 4, a second embodiment of the glazing strip 30 has recesses 32, 34 each for receiving a sheet of glass. The angle of the recess is 80 degrees. The adhesive strip 30 forms a 45 degree junction between the glass sheets.

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Referring to Figure 5 a third embodiment of the glazing strip 50 has recesses 52, 54 each for receiving a sheet of glass. The angle of the recess is 80 degrees. The adhesive strip so forms a 90 degree junction between the glass sheets.

The strip may be designed to permit junctions of different angles at any value between those shown in the Figures.

Referring to Figure 6, a fourth embodiment of the glazing strip 60 has recesses 62, 64, 66 each for receiving a sheet of glass thus forming a three-way junction.

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The glazing strip can have more than three recesses, for example four recesses for forming a 4-way junction.

CLAIMS

- 1. A composition for glazing joints comprising
- (a) a siloxane polymer having a molecular weight of from 300,000 to 700,000;
- 5 (b) a siloxane polymer having a molecular weight of from 10,000 to 100,000; the ratio of component A to component B being in the range of from 10 to 1, to 3 to 1; and
 - (c) a cross linking agent.
- 10 2. A composition as claimed in claim 1, wherein component (a) is a polymer containing vinyl groups.
 - 3. A composition as claimed in claim 1 or 2, wherein component (b) does not contain any vinyl groups.
 - 4. A composition as claimed in any preceding claim, wherein component (a) has a density of from 1.1 g cm⁻³ to 1.2 g cm⁻³
- 5. A composition as claimed in any preceding claim, wherein component (b) has a density of from s.g. 70 to 130.
 - 6. A composition as claimed in any preceding claim, wherein component (a) has a Shore hardness of from 10 to 95 °A.
- 25 7. A composition as claimed in any preceding claim, wherein component (a) has a Shore hardness of from 40 to 90 °A.
 - 8. A composition as claimed in any preceding claim, wherein component (b) has a viscosity of from 0.65 to 100,000 centistokes.

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- 9. A composition as claimed in any preceding claim, wherein component (b) has a viscosity of from 40,000 to 80,000 centistokes.
- 10. A composition as claimed in any preceding claim, wherein component (c) is a free radical initiator.
 - 11. A composition as claimed in any preceding claim, wherein component (c) is an organic peroxide.
- 10 12. A composition as claimed in any preceding claim, comprising from 80 to 90% component (a) by weight.
 - 13. A composition as claimed in any preceding claim, comprising about 86% component (a) by weight.
 - 14. A composition as claimed in any preceding claim, comprising from 5 to 20% component (b) by weight.
- 15. A composition as claimed in any preceding claim, comprising about 13% component (b) by weight.
 - 16. A composition as claimed in any preceding claim, comprising from 1 to 10% component (c) by weight.
- 25 17. A composition as claimed in any preceding claim, comprising about 5% component (c) by weight.
 - 18. A method of joining glazing sheets comprising
- (a) providing an adhesive strip shaped for receiving two or more glazing
 sheets, the surface of the strip being inherently adhesive;

- (b) inserting the strip between adjacent edges of at least a first and second sheet.
- 19. A method as claimed in claim 18, wherein the adhesive strip comprises a
- 5 compound as claimed in any of claims 1 to 17.

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FIGURE 1

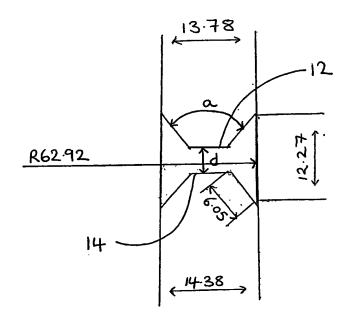
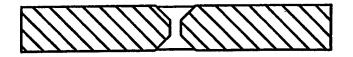


FIGURE 2



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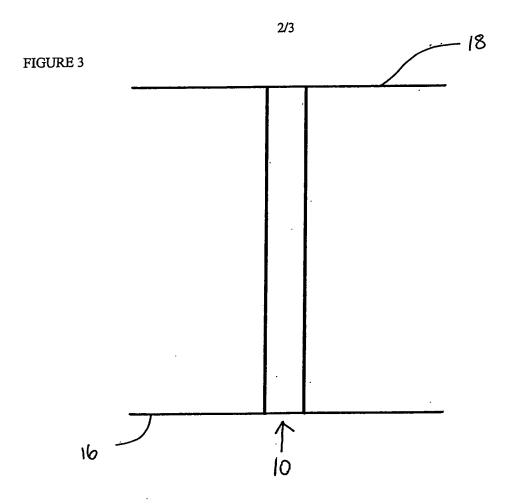
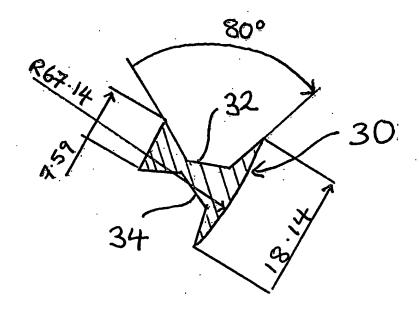


FIGURE 4



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FIGURE 5

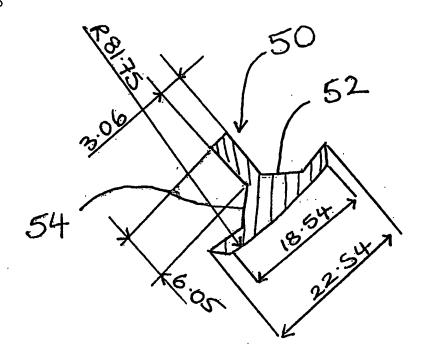
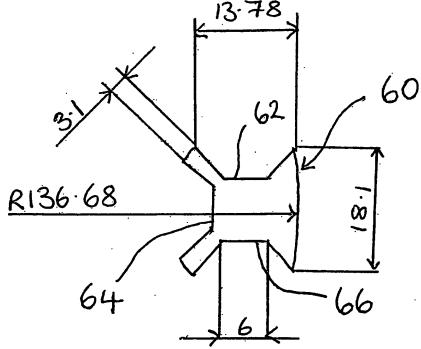


FIGURE 6



SUBSTITUTE SHEET (RULE 26)

INTERNATIONAL SEARCH REPORT

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A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 C09J183/04 E06B3/62 C03C27/10 According to International Patent Classification (IPC) or to both national classification and IPC B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) IPC 7 C09J E06B C03C C08L C08J Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the international search (name of data base and, where practical, search terms used) EPO-Internal, WPI Data, PAJ C. DOCUMENTS CONSIDERED TO BE RELEVANT Relevant to claim No. Citation of document, with indication, where appropriate, of the relevant passages GB 2 263 663 A (METABOWERKE KG) 1,18 4 August 1993 (1993-08-04) cited in the application claims 1,6-18 page 22, line 23 -page 23, line 18 Α US 5 584 957 A (SCHULTHEIS BERND ET AL) 1,18 17 December 1996 (1996-12-17) claims 1-11 column 1, line 8 - line 16 column 2, line 11 - line 18 column 4, line 64 -column 5, line 40 WO 94 01496 A (WINN RALPH ALASTAIR ; RILEY 1 Α NORMAN L (US); BRACKEN RONALD L (US); S) 20 January 1994 (1994-01-20) claims 1.3-7 -/--Y Further documents are listed in the continuation of box C. Patent family members are listed in annex. Special categories of cited documents: "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the *A* document defining the general state of the art which is not considered to be of particular relevance Invention "E" earlier document but published on or after the international filing date "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art. "O" document referring to an oral disclosure, use, exhibition or *P* document published prior to the international filing date but later than the priority date claimed "&" document member of the same patent family Date of the actual completion of the International search Date of mailing of the international search report 23/02/2004 16 February 2004 Name and mailing address of the ISA Authorized officer European Patent Ciffice, P.B. 5818 Patentitean 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016 Depijper, R

INTERNATIONAL SEARCH REPORT

PCT/GB 03/04467

	ation) DOCUMENTS CONSIDERED TO BE RELEVANT	 Bolovent to pint - Ma
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 99 03424 A (FRUCI VITTORIA) 28 January 1999 (1999-01-28) claims 1-24 page 3, line 31 - line 37 page 4, line 1 - line 35 page 13, line 29 -page 14, line 8	1
A	EP 1 201 705 A (DOW CORNING TORAY SILICONE) 2 May 2002 (2002-05-02) page 2, paragraph 2	1
A	gB 2 189 253 A (CENTRAL ELECTR GENERAT BOARD) 21 October 1987 (1987-10-21) claim 1 page 2, line 20 -page 3, line 25 page 5, line 15 - line 33	1

INTERNATIONAL SEARCH REPORT

PCT/GB 03/04467

Patent document cited in search report		Publication date		Patent family member(s)	<u></u>	Publication date
GB 2263663	A	04-08-1993	DE US	4202195 5332163		29-07-1993 26-07-1994
US 5584957	A	17-12-1996	DE	4327475		23-02-1995
	•		AT	191439		15-04-2000
			DE	59409262		11-05-2000
			EP	0639538		22-02-1995
			ES	2145790		16-07-2000
			JP	7102219	A 	18-04-1995
WO 9401496	Α	20-01-1994	AT	167220	Т	15-06-1998
			ΑU	666753	B2	22-02-1996
			ΑU	4544793	Α	31-01-1994
			CA	2139643	A1	20-01-1994
			DE		D1	16-07-1998
			DE		T2	29-10-1998
			DK	649447		29-03-1999
			EP	0649447		26-04-1995
			ES		T3	01-08-1998
			GR	3027770		30-11-1998
			JP	2802167		24-09-1998
			JP	7504937		01-06-1995
			MX	9304006		31-01-1995
			WO	9401496		20-01-1994
			US	5539020	A 	23-07-1996
WO 9903424	Α	28-01-1999	AU	8660798	Α	10-02-1999
			BR	9810775		19-09-2000
			WO	9903424	A1	28-01-1999
EP 1201705	A	02-05-2002	JP	2002129017	Δ	09-05-2002
Li 1201703	,.	02 00 2002	CA	2359767		25-04-2002
			EP	1201705		02-05-2002
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GB 2189253		21-10-1987	NONE			